

IN THE CLAIMS

Claim 1 (Original): A hollow fiber incorporating an immobilized biological substance, a porous fiber incorporating an immobilized biological substance, or a porous hollow fiber incorporating an immobilized biological substance, wherein the biological substance is directly immobilized on and/or in the fiber.

Claim 2 (Original): A fiber retaining a gel which incorporates an immobilized biological substance whereby the biological substance is immobilized on and/or in the fiber.

Claim 3 (Original): The fiber according to claim 2, which is a solid, hollow, porous or hollow porous fiber.

Claim 4 (Original): The fiber according to claim 3, which is a solid fiber, and wherein the gel incorporating an immobilized biological substance is retained on a surface of the fiber.

Claim 5 (Original): The fiber according to claim 3, which is a hollow fiber, and wherein the gel incorporating an immobilized biological substance is retained in a hollow part of the fiber.

Claim 6 (Original): The fiber according to claim 3, which is a porous fiber, and wherein the gel incorporating an immobilized biological substance is retained in the pore(s) of the fiber.

Claim 7 (Original): The fiber according to claim 3, which is a porous hollow fiber, and wherein the gel incorporating an immobilized biological substance is retained in a hollow part and the pore(s) of the fiber.

Claim 8 (Original): The fiber according to claim 1 or 2, wherein the biological substance is any one selected from a group consisting of the following substances (a) to (c):

- (a) nucleic acid, amino acid, sugar or lipid;
- (b) a polymer consisting of one or more kinds of ingredients from the substances stated in (a) above; and
- (c) a substance interacting with substances stated in (a) or (b) above.

Claim 9 (Original): The fiber according to claim 8, wherein the biological substance is nucleic acid.

Claim 10 (Original): The fiber according to claim 2, also having a pigment retained on and/or in the fiber by means of the gel.

Claim 11 (Original): A fiber alignment having a bundle of the fibers stated in any one of claims 1 to 10.

Claim 12 (Original): The fiber alignment according to claim 11, wherein the fibers are regularly arranged.

Claim 13 (Original): The fiber alignment according to claim 11, wherein the bundle of the fibers comprises 100 or more fibers per cross-sectional cm^2 .

Claim 14 (Original): The fiber alignment according to claim 11, wherein the type of biological substance on each fiber is different in respect of some or all of the fibers.

Claim 15 (Original): A slice of the fiber of claim 11, which intersects the fiber axis of the fiber alignment according to claim 11.

Claim 16 (Original): The slice according to claim 15 comprising fiber units and coordinates reference points therefor.

Claim 17 (Original): The slice according to claim 16, wherein the coordinate reference points are two or more marker fiber units therein.

Claim 18 (Original): The slice according to claim 17, wherein the marker fiber units are stained.

Claim 19 (Original): The slice according to claim 16, wherein the coordinates for a fiber unit are determined based on the coordinate reference points.

Claim 20 (Original): A method for producing the slice according to claim 16 having coordinates for each fiber unit thereof, the method comprising the steps of:

- (a) cutting sequentially a fiber alignment obtained by binding and immobilizing fibers, to obtain a series of fiber alignment slices S(1), S(2), ... S(h), ... S(m);

- (b) selecting any given slice $S(h)$ from m number of slices and determining two-dimensional coordinates for each fiber unit contained in said slice $S(h)$ based on the coordinate reference pointss in said slice $S(h)$;
- (c) determining the two-dimensional coordinates of each fiber unit contained in slice $S(i)$ located close to said slice $S(h)$ based on the coordinate data of slice $S(h)$ obtained in step (b) and the coordinate reference points in said slice $S(i)$; and
- (d) repeating steps (b) and (c) to determine the two-dimensional coordinates of each fiber unit in said fiber alignment slice.

Claim 21 (Currently Amended): A method for determining the position of each fiber unit in the slice according to claim 16, the method comprising the steps of:

- (a) cutting sequentially a fiber alignment obtained by binding and immobilizing fibers, to obtain a series of fiber alignment slices $S(1)$, $S(2)$, ... $S(h)$, ... $S(m)$;
- (b) selecting any given slice $S(h)$ from m number of slices, and determining two-dimensional coordinates for each fiber unit contained in said slice $S(h)$, based on the coordinate reference ~~pointss~~ points in said slice $S(h)$;
- (c) determining the two-dimensional coordinates of each fiber unit contained in slice $S(i)$ located close to said slice $S(h)$, based on the coordinate data of slice $S(h)$ obtained in step (b) and the coordinate reference points in said slice $S(i)$; and
- (d) repeating steps (b) and (c) to determine the two-dimensional coordinates of each fiber unit in said fiber alignment slice.

Claim 22 (Original): A computer-readable recording medium on which the coordinate data of each fiber unit in the slice according to claim 16 is recorded.

Claim 23 (Original): A set for sample detection, comprising slices according to claim 16 and the recording medium according to claim 22.

Claim 24 (Original): A method for producing the slice according to claim 15, which comprises: binding a plurality of hollow fibers to make an alignment; introducing a biological substance into the inner wall and/or hollow part(s) of each hollow fiber constituting said alignment and immobilizing the substance therein; and slicing the said alignment in a direction intersecting with the fiber axis.

Claim 25 (Original): A method for producing the slice according to claim 15, which comprises: binding a plurality of porous hollow fibers to make an alignment; introducing a biological substance into the inner wall, hollow and/or porous part(s) of each porous hollow fiber constituting said alignment and immobilizing the substance therein; and slicing the said alignment in a direction intersecting with the fiber axis.

Claim 26 (Original): The method according to claim 24, wherein the immobilization of a biological substance in the inner wall and/or hollow part(s) of each hollow fiber constituting an alignment is carried out by immersing the extended tip of each hollow fiber constituting said alignment into a solution containing a biological substance, and introducing said solution into the hollow part of each hollow fiber constituting said alignment.

Claim 27 (Original): The method according to claim 25, wherein the immobilization of a biological substance in the inner wall, hollow and/or porous part(s) of each porous hollow fiber constituting an alignment is carried out by immersing the extended tip of each porous hollow fiber constituting said alignment into a solution containing a biological

substance, and introducing said solution into the hollow and/or porous part(s) of each porous hollow fiber constituting said alignment.

Claim 28 (Original): A method for producing a fiber alignment, which comprises applying tension to a fiber bundle arranged in accordance with a sequence pattern of interest, and immobilizing said fiber bundle by filling resin among fibers of said fiber bundle to make a fiber alignment.

Claim 29 (Original): The production method according to claim 28, wherein the sequence of a fiber bundle is formed by the steps of:

- (a) passing fibers through a plurality of jigs having pores of the same pattern as a sequence pattern of interest; and
- (b) widening the intervals between said jigs.

Claim 30 (Original): The production method according to claim 29, wherein the jigs are support lines constituting networks obtained by longitudinal and transverse lines, or a perforated board.

Claim 31 (Original): A method for treating the inner wall part of a hollow fiber, which comprises applying a gel forming monomer (a) solution on the inner wall of a hollow fiber, and then forming gel on the inner wall of said hollow fiber by polymerization of said monomers.

Claim 32 (Original): The method according to claim 31, wherein the inner wall is porous.

Claim 33 (Original): The method according to claim 31, wherein monomer (a) is an amphipathic monomer.

Claim 34 (Original): A method for filling the hollow part of a hollow fiber with gel, which comprises filling a gel forming monomer (b) solution in the hollow part of a hollow fiber treated by any one of the methods according to claims 31 to 33, and forming gel in the hollow part by polymerization of said monomers.

Claim 35 (Original): The method according to claim 34, wherein the main ingredient of monomer (b) is acrylamide.

Claim 36 (Original): A method for producing a gel-filled fiber, which comprises filling the hollow part of a hollow fiber treated by any one of the methods according to claims 31 to 33 with a gel forming monomer (b) solution, and forming gel in the hollow part by polymerization of said monomers.

Claim 37 (Original): The production method according to claim 36, wherein the main ingredient of monomer (b) is acrylamide.

Claim 38 (Original): A polymer gel incorporating immobilized nucleic acid, wherein modified nucleic acid is bound and immobilized thereon by means of a glycidyl group.

Claim 39 (Original): The polymer gel according to claim 38, wherein the modified nucleic acid has an aminated terminus.

Claim 40 (Original): The polymer gel according to claim 38, wherein the polymer gel is a copolymer gel consisting of glycidyl(meta)acrylate, a polymerized monomer and a cross-linker.

Claim 41 (Original): The polymer gel according to claim 40, wherein the polymerized monomer is acrylamide.

Claim 42 (Original): A method for producing the polymer gel according to claim 38, which comprises reacting glycidyl(meta)acrylate with a modified nucleic acid and then adding a polymerized monomer and a cross-linker to the obtained reaction product to polymerize them.

Claim 43 (Original): A method for producing the polymer gel according to claim 38, which comprises reacting modified nucleic acid with a copolymer gel consisting of glycidyl(meta)acrylate, a polymerized monomer and a cross-linker.

Claim 44 (Original): The production method according to claim 42 or 43, wherein the modified nucleic acid has an aminated terminus.

Claim 45 (Original): The production method according to claim 42 or 43, wherein the polymerized monomer is acrylamide.

Claim 46 (Original): A polymer gel comprising a nucleic acid ingredient, a polyvalent amine ingredient and at least two or more polymerized monomer ingredients.

Claim 47 (Original): The polymer gel according to claim 46, wherein at least one of polymerized monomer ingredients is a polymerized monomer having a glycidyl group.

Claim 48 (Original): The polymer gel according to claim 47, wherein the polymerized monomer having a glycidyl group is glycidyl(meta)acrylate.

Claim 49 (Original): The polymer gel according to claim 46, wherein the nucleic acid ingredient has an aminated terminus.

Claim 50 (Original): A method for producing the polymer gel according to claim 46, which comprises polymerizing a solution comprising a nucleic acid ingredient, a polyvalent amine ingredient and at least two or more polymerized monomer ingredients.

Claim 51 (Original): A method for producing the polymer gel according to claim 46, which comprises polymerizing a solution comprising a nucleic acid ingredient and at least two or more polymerized monomer ingredients, and cross-linking the obtained polymer with a polyvalent amine ingredient.

Claim 52 (Original): A method for detecting a sample which comprises using the slice according to claim 15, the slice having as a probe a biological substance attached to a carrier, wherein said method comprises bringing the sample into contact with said slice by a method other than natural diffusion to form a hybrid, and removing from said slice samples which do not bind to the biological substance probe.

Claim 53 (Original): The detection method according to claim 52, wherein the sample is brought into contact with the slice by applying a voltage across said slice.

Claim 54 (Original): The detection method according to claim 52, wherein a water-absorbing substance is located on one side of said slice thereby bringing a sample located on the opposite side into contact with the slice.

Claim 55 (Original): The detection method according to claim 52, wherein the biological substance is nucleic acid.

Claim 56 (Original): The detection method according to claim 52, wherein the sample is labeled by fluorescence.

Claim 57 (Original): The detection method according to claim 52, wherein the carrier is a soluble polymer gel.

Claim 58 (Original): The detection method according to claim 57, wherein the main ingredient of the soluble polymer gel is polyacrylamide.

Claim 59 (Original): The detection method according to claim 52, wherein the carrier is retained in the hollow part of a hollow fiber.